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<110> Stanton, Jr., Vincent P.

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<211> 3951

<212> DNA

<213> Homo sapiens

<220>

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<222> 577, 638, 1708, 3730, 3925

<223> n = a or g

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 <213> Homo sapiens

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 <222> 175, 1067
 <223> n = g or a

<221> misc_feature
 <222> 341
 <223> n = c or g

<221> misc_feature
 <222> 791, 1997, 2618, 2653
 <223> n = t or c

<221> misc_feature
 <222> 1337
 <223> n = c or a

<221> misc_feature
 <222> 2107
 <223> nucleotide in position 2107 is g, or absent

<221> misc_feature
 <222> 2583
 <223> n = t or g

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<211> 3772
<212> DNA
<213> Homo sapiens

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<223> n = a or g

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<221> misc_feature
<222> 498
<223> n = c or t

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<221> misc_feature
<222> 579, 599
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<212> DNA
<213> Homo sapiens

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<222> 1066
<223> n = t or c
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<221> misc_feature

<222> 1136

<223> n = a or g

<221> misc_feature

<222> 1497

<223> n = t or a

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<211> 1187

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<220>

<221> misc_feature

<222> 276, 321, 534, 656

<223> n = c or t

<221> misc_feature

<222> 452, 640

<223> n = a or g

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<222> 492, 625

<223> n = c or a

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<222> 458

<223> nucleotide in position 458 is c, or absent

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 15503, 15590, 15840, 16149
 <223> n = a or g

<221> misc_feature
 <222> 732, 1379, 1590, 2488, 3212, 5006, 11238, 11422, 11686,
 12598, 13171, 13645, 13782, 13806, 13813, 14586, 14788,
 15042, 15546, 15770
 <223> n = c or t

<221> misc_feature
 <222> 1322, 1688
 <223> n = c or g

<221> misc_feature
 <222> 2594, 11293, 16199, 16203
 <223> n = g or t

<221> misc_feature
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 <223> n = a or t

099333-09401

<221> misc_feature

<222> 14547

<223> nucleotide in position 14547 is t, or absent

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<222> 1045

<223> nucleotide at position 1045 is c, or absent

<221> misc_feature

<222> 1046

<223> nucleotide at position 1046 is t, or absent

0996333.09401

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 <222> 2636, 5287
 <223> n = c or g

<221> misc_feature
 <222> 3118
 <223> n = g or t

<221> misc_feature
 <222> 3257, 4053
 <223> n = a or c

<221> misc_feature
 <222> 5440
 <223> n = t or a

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<211> 2158
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<213> Homo sapiens

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<223> n = c or t

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0996333-09404

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 <212> DNA
 <213> Homo sapiens

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 <222> 263, 1037, 1139, 1955, 2017, 2037, 2189, 2309
 <223> n = a or g

<221> misc_feature
 <222> 266
 <223> n = g or t

<221> misc_feature
 <222> 527
 <223> n = c or g

<221> misc_feature
 <222> 1217, 1647, 2282
 <223> n = c or t

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<221> misc_feature

<222> 464

<223> n = g or t

<221> misc_feature

<222> 120, 519, 668, 1059, 1308

<223> n = c or t

<221> misc_feature

<222> 1289

<223> n = c or a

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<213> Homo sapiens

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<222> 575, 648

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<221> misc_feature

<222> 771

<223> n = g or c

<221> misc_feature

<222> 883

<223> n = g or a

<221> misc_feature

<222> 942

<223> nucleotide at position 942 is c, or absent

<221> misc_feature

<222> 1052

<223> n = a or c

<400> 16

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